

ULTRA- chloroxylenol 0.3% liquid

Aldi

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug facts Radiance 956.002/956AE

Active ingredient

Chloroxylenol 0.3%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only: hands only

When using this product

- do not get into eyes. If contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

water, sodium C14-16 olefin sulfonate, lauramine oxide, sodium laureth sulfate, sodium lauryl sulfate, sodium xylenesulfonate, sodium chloride, alcohol denat., fragrance, phenoxyethanol, citric acid, methylisothiazolinone, tetrasodium EDTA, yellow 5, blue 1

WARNING

NOT FOR USE IN DISHWASHERS.

DO NOT ADD BLEACH

Phosphate Free Contains Surfactants

Not tested on animals

ANTIBACTERIAL

HAND SOAP

GREEN APPLE SCENT

Ultra Concentrated

Dishwashing Liquid

DIST. & SOLD EXCLUSIVELY BY:

ALDI, BATAVIA, IL 60510

This product is not manufactured or distributed by Procter & Gamble, distributor of Dawn Ultra Antibacterial Hand Soap Apple Blossom Scent

principal display panel

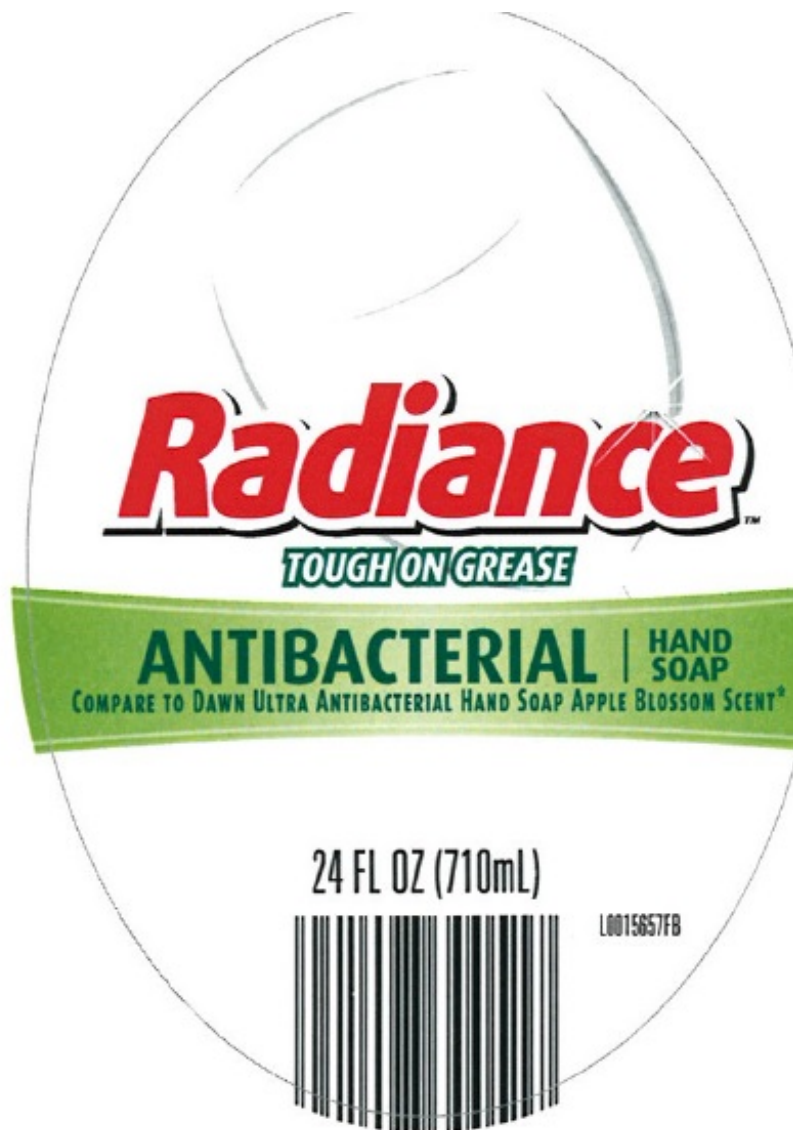
Radiance

TOUCH ON GREASE

ANTIBACTERIAL | HAND SOAP

COMPARE TO DAWN ULTRA ANTIBACTERIAL HAND SOAP APPLE BLOSSOM SCENT

24 FL OZ (710 mL)



ULTRA

chloroxylenol 0.3% liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64024-956
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	3.09 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	

SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALCOHOL (UNII: 3K9958V90M)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
EDETATE SODIUM (UNII: MP1J8420LU)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64024-956-50	710 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/17/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/17/2019	

Labeler - Aldi (944259522)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(64024-956)

Revised: 4/2021

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